Please amend page 28, line 1 as follows:

Claims What is claimed is:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Original) A compound of the general formula I:

$$V-L-Z$$

Formula I

or pharmaceutically acceptable salt thereof, wherein V is a non-peotidic vector having affinity for the Angiotensin II receptor L is a bond, a spacer or a linker moiety and Z represents a moiety detectable in an *in vivo* imaging procedure of a human or animal body.

- 2. (Original) A compound according to claim 1 wherein V is Losartan, Valsartan, Candesartan, Eprosartan or derivatives thereof.
- 3. (Currently amended) A compound as claimed in any of the previous claims claim 1, where Z is a chelating agent of Formula II carrying an imageable moiety M

where:

each R^1 , R^2 , R^3 and R^4 is independently an R group; each R group is independently H or C_{1-10} alkyl, C_{3-10} alkylaryl, C_{2-10} alkoxyalkyl, C_{1-10} hydroxyalkyl, C_{1-10} alkylamine, C_{1-10} fluoroalkyl, or 2 or more R groups, together with the atoms to which they are attached form a carbocyclic, heterocyclic, saturated or unsaturated ring.

4. (Currently amended) A compound as claimed in any of the previous claims claim 1, where Z is a chelating agent of formula e carrying an imageable moiety M

- 5. (Currently amended) A compound as claimed in any of the previous claims claim 1, wherein Z comprises an imaging moiety wherein the imaging moiety comprises metal radionuclides, paramagnetic metal ions, fluorescent metal ions, choromophores, heavy metal ions or cluster ions.
- 6. (Currently amended) A compound as claimed in elaims 3–5 claim 3, wherein the imaging moiety comprises ⁹⁰Y, ^{99m}Tc, ¹¹¹In, ⁴⁷Sc, ⁶⁷Ga, ⁵¹Cr, ^{177m}Sn, ⁶⁷Cu, ¹⁶⁷Tm, ⁹⁷Ru, ¹⁸⁸Re, ¹⁷⁷Lu, ¹⁹⁹Au, ²⁰³Pb, ¹⁴¹Ce or ¹⁸F.
- 7. (Original) A pharmaceutical composition comprising an effective amount of a compound of general Formula (I) or a salt thereof, together with one or more pharmaceutically acceptable adjuvants, excipients or diluents for use in enhancing image contrast in *in vivo* imaging or for treatment of a disease.

- 8. (Currently amended) Use of a compound as claimed in any one of claims 1 to 6 claim 1, in the preparation of a contrast medium for use in a method of diagnosis involving administering said contrast medium to a human or animal body and generating an image of at least part of said body.
- 9. (Currently amended) A method of generating images of a human or animal body involving administering a contrast agent to said body, and generating an image of at least a part of said body to which said contrast agent has distributed, characterised in that said contrast agent comprises a compound as claimed in any one of claims 1 to 6 claim 1.
- 10. (Currently amended) A method of generating enhanced images of a human or animal body previously administered with a contrast agent composition comprising a compound as claimed in elaims 1 to 6claim 1, which method comprises generating an image of at least part of said body.